IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

First Applicant:	John A. Hamilton
Serial No.:	09/851,230
Filing Date:	May 8, 2001
Group Art Unit:	1644
Examiner:	Belyavskyi, Michail A.
For:	A Method for the Treatment and Prophylaxis of Inflammatory Conditions

DECLARATION UNDER 37 C.F.R. § 1.131

We, John A. Hamilton and Gary P. Anderson, declare that:

- 1. We are joint inventors of the subject matter of the above-identified patent application (the "230 application"); the contents of the application are well known to us and we have read the Office Actions dated August 10, 2005 and March 28, 2006 as well as U.S. Patent Application Publication No. US2000-0141994 to Devalaraja, *et al.* (the "994 Publication").
- 2. We carried out, or caused to be carried out at our direction and under our supervision, certain experimental work which demonstrates that the invention of the '230 application was completed prior to March 20, 2000.
- 3. A red number has been added in the lower left hand corner of the attached notebook pages to facilitate reference thereto. The dates have also been redacted. No other changes have been made to the original documents.
- 4. The experimental work described herein occurred prior to March 20, 2000.

 Scientific data evidencing the conception and reduction to practice of the present invention from

the contents of attached Exhibit 1. The data come from our scientific laboratory notebooks and demonstrate that antibodies specific to GM-CSF are capable of ameliorating the effects of inflammation in a mouse by inhibiting or antagonizing the effects of GM-CSF on cells of the monocyte/macrophage lineage.

- 5. We observed that mice with a knockout mutation in GM-CSF (GM-CSF -/-) demonstrated a significantly reduced incidence of arthritis compared to wild-type mice (GM-CSF +/+) or heterozygous mice (GM-CSF -/+). This indicated that inhibiting GM-CSF could be a means of treating or preventing inflammation. Based thereon, we conducted experiments to use antibodies to specifically inhibit GM-CSF.
- 6. DBA\1 mice were injected with type II collagen (CII) in Freund's Complete Adjuvant (FCA) to induce arthritis (Exhibit 1, top of p.1). After appearance of arthritis, these mice (10 per group) were either treated with 22E9, a neutralizing monoclonal antibody specific for GM-CSF or an isotype control (Exhibit 1, middle of p.1). The clinical score was monitored daily to ascertain the level of inflammatory arthritis present within the two groups of animals. As seen in Figure 1 (Exhibit 1, p. 3), animals which were treated with the 22E9 monoclonal antibody specific for GM-CSF developed significantly lower levels of disease than those treated with the isotype control.
- 7. Attached Exhibit 2 contains data from our scientific laboratory notebooks that demonstrate that antibodies specific for M-CSF are capable of ameliorating inflammatory arthritis in a mouse by inhibiting or antagonizing the effect of M-CSF on cells of the monocyte/macrophage lineage.
- 8. We observed that mice with a mutation rendering M-CSF inactive demonstrated significantly reduced arthritis compared to wild-type mice. This indicated that inhibiting M-CSF

could be a means of treating or preventing inflammation. Based thereon, we conducted experiments to use antibodies to specifically inhibit M-CSF.

- 9. DBA/1 mice were injected with CII in FCA to induce arthritis as described on page 1 of Exhibit 2. At day 27, mice exhibiting signs of arthritis were divided into 2 groups of 10 and injected with either 5AI, a monoclonal antibody specific for M-CSF, or an isotype control antibody, DX48. The clinical score was monitored daily to ascertain the level of inflammatory arthritis present within the two groups of animals. As seen in Figure 2 (Exhibit 2, page 3), animals treated with the 5AI monoclonal antibody specific for M-CSF developed significantly lower levels of disease than those treated with the isotype control.
- 10. Accordingly, prior to March 20, 2000, we demonstrated that antibodies specific for GM-CSF or M-CSF are capable of ameliorating the effects of inflammation in a subject by inhibiting or antagonizing the effects of GM-CSF or M-CSF on cells of the monocyte/macrophage lineage.
- and that all statements made on information and belief are believed to be true; and further, that these statements were made with the knowledge the willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the application or any patent issuing thereon.

Date: 22 John A. Hamilton

Date: 22" June, 2006 Gary Anderson

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EXHIBIT 1

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	Effect of stole on lake onset mice us Ratify. (CIASE NO)
	Effect of 3466 on lake onset mile us Ratify. (CIASE Not. 22E9. (d28x) (d27-2) 7-11 whos OR: Strain: OBA/1 Lac-J
	#1-41 (II:FCA i.d. 100 pt MJR +7 died.
	#1-41 CT: Flxid 100/21 MJR #26 sick odied
	Mice scared daily - don't need to do astually
	alice need to do as mice come up early
	Score all Mile. HALLE up by now overit melucled in
	sample group & controll group mice, balance the
•	Maria Waller Maria Committee Committ
	Gave 22E9-0.3mg (110 pl) & 2.65 mg/ml purific
	Gave 22E9-0.3mg (110 pl) @ 2.65 mg/ml purific i.e. in PBS. From Cell Max beginning Robbs -s/www.s. podled & who year purified & dicolysed 1. To PBS. indooring = 210pg/ml
	Rut Iqq- 4s for (1457
· .	Treatment-0.3 mg of Abs i.p. For 10 consecutive clarge from of 27 post immunisation. Swollen tops counted
	22E9 *1, *9 *14 *16 *18 *21 *22 *30 *31 *38
	Ratiq *15 *20 "24 "27 +33 +34 *35 *36 #37 #40
	Mile killed on d15 & post initial Ab Frecetment Blood tak Rear limbs 8 splans Rixed
	(NB). Next 2269 CIA expt - need to take out
	the lungs of mice given 2159 and those given control

	CIA Practicia
P.	actising collage id injections in DBA/1 mice and including
a	thritis in these mice.
•	F12, WKS,
#	2-6 and 8-17 (15 mile) CIL FCA id. 10gal Emma (2-5216) Natalie (#6,8-10,17) Fiona (#11-15) #1 and #7 died.
4	# 2-5 8-16 (13 mice) CII: F(A id 100 nL Emma (#2-5) #### Natalie (#8-10, #16) Frona (#11-15) # 6 died , # 17 was missing !
	# 6 died # 17 was missing
-	
<u> </u>	Score mice dially - come had come up already, need to score earlier next time.
1	
	Kill DBA # 10 and # 11. (ie # 10 Scored 10 #11 = 0) Collect blood, and limbs. Do a paritoneal louvage to onah cells by FACS.
	Kill - miles collect blood by heart ouncture
	Kill remaining mice, collect blood by heart puncture,
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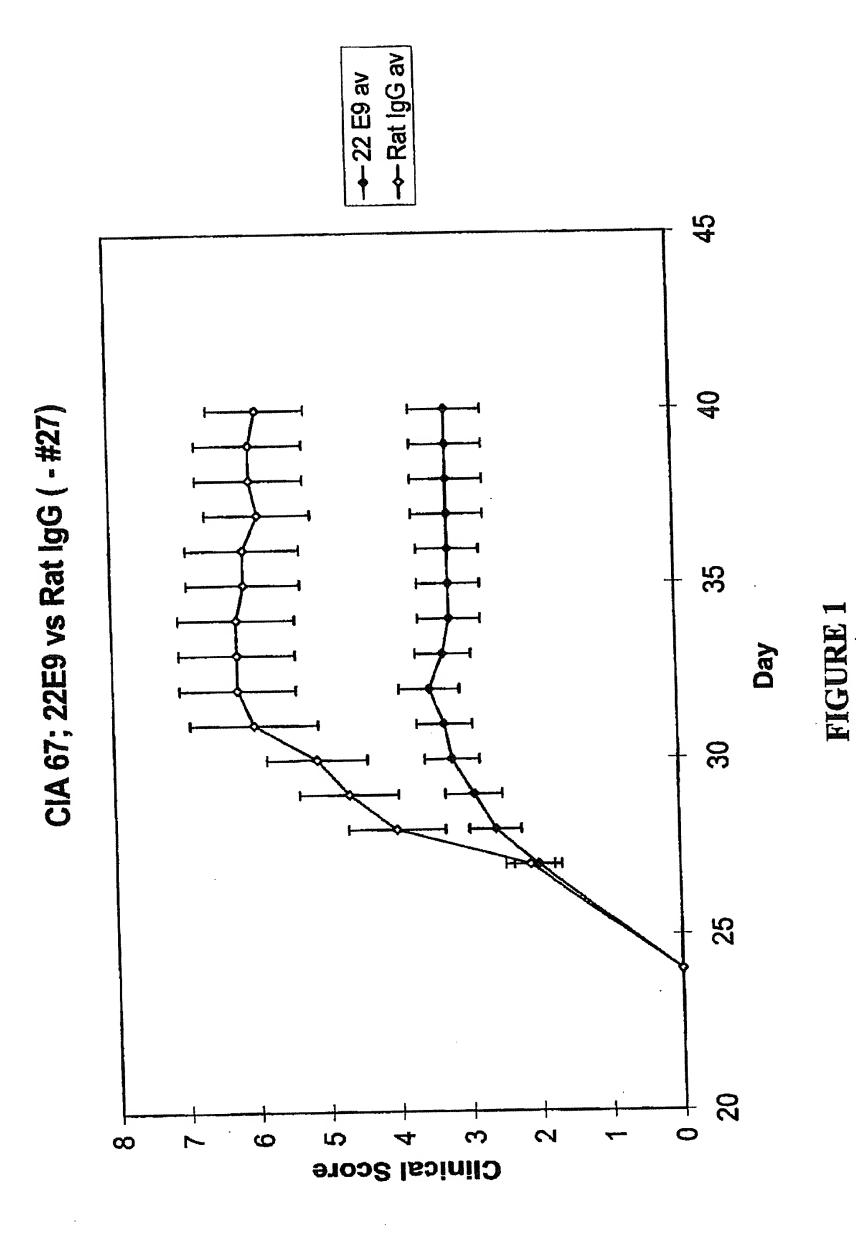


EXHIBIT 2

CIAGS - 5AT on established# | 41 000 DBA/1 agget 9-13 soles now

- injected = cII | CFA as par usual.

IKC injected # 1-20

MR is # 21-30

JD is # 31-40

AM mice impeded & CIT boost in CFA (11:1).

Newly the mice arrighed to one of two injection groups.

(a) 5AI (300µg; i.p.) ... 10 mice.

(b) DX4-8 (300µg; i.p.) ... 10 mice.

for 10 consecutive doinly injections.

(0.2 ml of 1.5 mg/ml in PBS)

Much worse on the Med on they continued their

Mile were marriaded Med spleans & all lunder semi-order

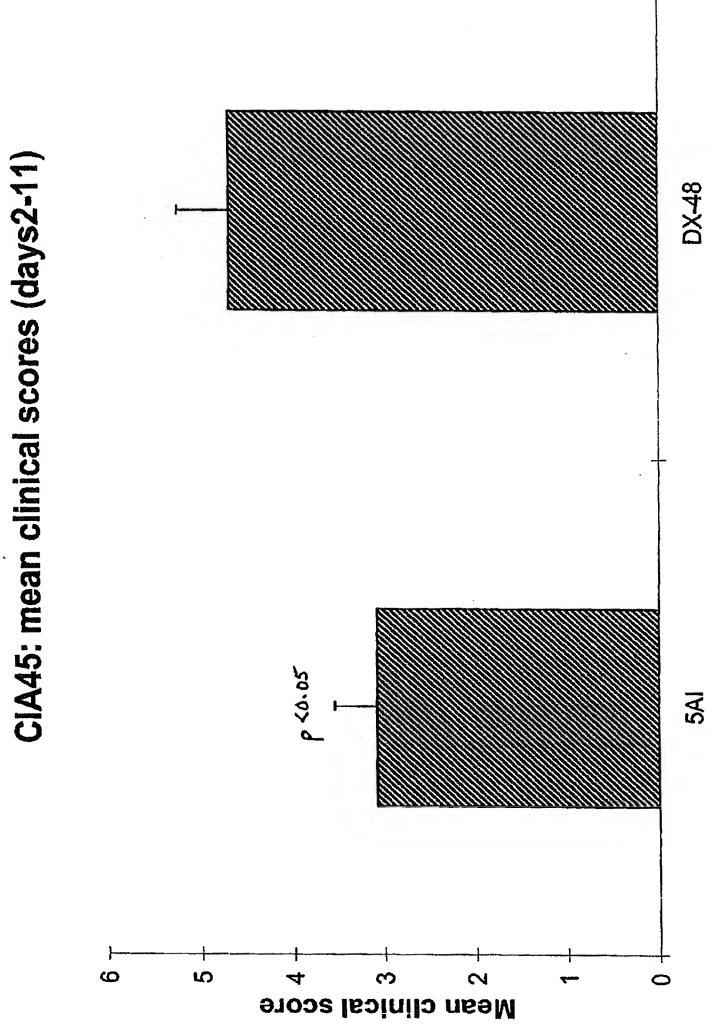
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CIA-45. Day of onzet.

ommitted - 26,26,26,26,26,26,26,26,24,26,24,

 $5AI - 27, 28, 28, 27 = 29, 29, 30, 29, 27, 28.5 \pm 0.4$ 28.5 ± 0.4 28.4 ± 0.4 28.8 ± 0.6

13 12 9 CIA45: clinical scores O Days of arthritis AD TREMTMENT S → DX-48 * P <0.05 --- 5AI က ~ Citting... က ~ 9 Clinical score



Antibody treatment

CIA45.XLS

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